

K242776 LimFlow V-CeiverFeb 14, 2025
154 days to decisionK242776 · Product code: **MMX** · CardiovascularSource: <https://www.510kdatabase.net/k242776/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Device, Percutaneous Retrieval (MMX)
Date received	Sep 13, 2024
Decision date	Feb 14, 2025
Days to decision	154 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

APPLICANT

Company	LimFlow, Inc.
Location	San Jose, CA, US
Contact	Tosan Eweka
510(k) history	6 submissions · 6 cleared · 2022-2026

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k242776/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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